Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	1	("620 44 24").PN.	USPAT	OR	OFF	2005/07/18 07:15
L2	1	("20020137966").PN.	US-PGPUB	OR	OFF	2005/07/18 07:15
L3	1	("20030055279").PN.	US-PGPUB	OR	OFF	2005/07/18 07:16
L4	1	("20030109745").PN.	US-PGPUB	OR	OFF	2005/07/18 07:17
L5	1	("20030125578").PN.	US-PGPUB	OR	OFF	2005/07/18 07:30
L6	579	560/106 OR 560/240	USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2005/07/18 07:30
L7	292	L6 AND (ANHYDRIDE OR ANHYDRIDES) AND (ETHER OR ETHERS)	USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2005/07/18 07:31
L8	231	L7 AND CATALYST	USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2005/07/18 07:34
L9	16	L8 AND (POLYOXOMETALATE OR HETEROPOLYACIDS OR HETEROPOLYACID OR SILICOMOLYBDIC OR SILICOTUNGSTIC OR SILICOTUNGSTATE OR PHOSPHOTUNGSTATE OR PHOSPHOTUNGSTATE OR PHOSPHOVANADATE OR PHOSPHOVANADATE OR PHOSPHOVANADIC)	USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2005/07/18 07:36

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Ikeuchi, Yuteaka; Taguchi, Takeo; Hanzawa, Yuji
Sankyo Co. Ltd., Kanagawa, 254-8560, Japan
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Sumitomo Chemical Company, Ltd., Japan
U.S., 5 pp., Cont.-in-part of U.S. Ser. No. 637,904, abandoned.
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Department of Chemistry, Mazandaran University, Baboolsar, 407415/453,
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Bereich Org. Grundst., Zentralinst. Org. Chem., Lelpzig, O-7050, Germany
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Labranal Prikladnoi Khimii (Sankt-Peterburg, Russian Federation) (1990), 63(6), 1376-83 19740805 Short synthesis of C-aryl glucopyranosides of the papulacandin type Schmidt, Richard R.; Frick, Wendelin Fak. Chem., Univ. Konstanz, Konstanz, D-7750, Fed. Rep. Ger. Tetrahedron (1988), 44(23), 7163-9 CODEN: TETRAB; ISSN: 0040-4020 APPLICATION NO. ANSWER 10 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN 1976:446771 CAPLUS 85:46771 COPYRIGHT 2005 ACS on STN 19760330 19740805 DATE Pyrrolobenzodiazepines Wright, William Blythe, Jr. American Cyanamid Co., USA U.S., 10 pp. KIND A A ANSWER 8 OF 10 CAPLUS 1991:23247 CAPLUS 114:23247 English CASREACT 111:58184 CASREACT 114:23247 US 3947408 US 1974-494657 PATENT NO. English Journal Journal Patent LA Enc E A SCS SCATURAGE SEA 188 SE PT 片

Benzyl acetate from dibenzyl ether

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ANSWER 7 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN High-temperature and -pressure reaction of (PhCH2)20 with Ac20 gave PhCH2OAc. The mechanism and optimization conditions of the reaction are discussed.

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Experimental and calculated property data are now available. For moi information enter HELP PROP at an arrow prompt in the file or refer to the file summary abset on the web at: to the file armary abset on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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Sulfamic acid as a cost-effective catalyst instead of metal-containing ANSWER 1 OF 11 CAPLUS COPYRIGHT 2005 ACS on STN 2004:641912 CAPLUS 141:313882 T P A I

acids for acetolysis of cyclic ethers Wang, Bo; Gu, Yanlong; Gong, Weizhong; Kang, Yuru; Yang, Liming; Jishuan Ŋ

State Key Laboratory for Oxo Synthesis and Selective Oxidation, Lanzinstitute of Chamical Physics, Chinese Academy of Sciences, Lanzhou, 730000, Peop. Rep. China Tettahedron Letters (2004), 45(35), 6599-6602 CODEN: TELEAY; ISSN: 0040-4039 ន

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Elsevier

FOR THIS RECORD CASREACT 141:313882 T 10 THERE ARE 10 CITED REFERENCES AVAILABLE F ALL CITATIONS AVAILABLE IN THE RE FORMAT ž RSEAR

CAPLUS COPYRIGHT 2005 ACS on STN ANSWER 2 OF 11 CAP 2004:247001 CAPLUS HARI.

Esterification process and catalysts for the preparation of benzyl carboxylate esters from dibenzyl ethers and/or alcohols and carboxylic acid anhydrides

Ooms, Pieter; Schenke, Bernd-Ulrich Bayer Chemicals A.-G., Germany Eur. Pat. Appl., 11 pp. GODEN: EPXXDW S PA

German Patent DT Pat LA Ger FAN.CNT

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CASREACT 140:270626; MARPAT 140:270626 2002-10243200 PRAI OS ANSWER 3 OF 11 CAPLUS COPYRIGHT 2005 ACS on STN 2002:185052 CAPLUS

136:233896

Manufacture of carboxylic acid benzyl esters from dibenzyl ethers Ooms, Pieter, Schenke, Bernd-Ulrich NI SE SE S

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| Bayer Aktiengesellschaft, Germany PCT Int. Appl., 19 pp. | | | DATE | 48285 | SEST | NL, SE, MC
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| Bayer Aktiengesellschaft, Germany CODEN: PIXXD2 Patent CODEN: PIXXD2 Patent CODEN: PIXXD2 Patent COC CC, CC, CC, CB, DK, DM, DK, DM, DC, CC, CC, CC, CB, DK, DM, DM, CC, DK, DM, DC, DK, DK, DK, DK, DK, DK, DK, DK, DK, DK | - | | APPLICATION NO. | WO 2001-EP9677 BB, BG, BR, BY, EC, EE, ES, FI, KE, KG, KP, KR, MN, MW, MX, MZ, SL, TJ, TM, TR, | SZ, TZ, UG, ZW,
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Sulfonic-carboxylic anhydrides were prepared and used for the cleavage of ethers. Thus acetyl p-toluenesulfonate (I) was prepared (97.5%) by the reaction of p-toluenesulfonic acid and excess AcCl. A mixture of crude I, MeCN, and excess BuZO was refluxed to give 50% Bu tosylate and some BuOAc. Iso-PrZO, (phCHZ) 20 (to 50% PhCHZOAc and 30% benzyl tosylate) if IHF [to 73% AcO(CHZ)403ScGH4Mep], 2-methyltetrahydrofuran [to 95% AcO(CHZ)20MeoSSCGH4Me-p] and dioxane [to 87% AcO(CHZ)203ScGH4Me-p] and dioxane [to 87% AcO(CHZ)20GHZOGH4Me-p] used similarly. ANSWER 11 OF 11 CAPLUS COPYRIGHT 2005 ACS on STN L18 AB

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ANSWER 7 OF 11 CAPLUS COPYRIGHT 2005 ACS on STN A process for the preparation of carboxylate esters from cyclic or acyclic ethers comprises treatment with aliphatic or aromatic carboxylic anhydrides in

3:1-1:1 ratio at 450-630 K at 0.1-20 mPA in a reaction zone. The process produces esters free of catalysts. PMGHZOMe (0.3 mol) and Ac20 (0.3 mol) were contacted in a reaction zone at 573 K and 6.5 MPa initial hydrogen pressure for 20 min; the selectivity toward formation of McO2CHZPh was 42% and toward formation of McO2Me 31%; conversion was 50%. Heating PhCHZOMe (0.3 mol) and Ac20 (0.3 mol) at reflux for 23 h gave a conversion of 2.1% and a selectivity toward McO2CHZPh and McCOZMe of 10% and 8%,

NSWER 8 OF 11 CAPLUS COPYRIGHT 2005 ACS on STN process for the preparation of MeCO2CH2Ph from a hyproduct of perfume ANSWER 8 OF 11 L18 ANSWER EAB AB A process

comparises the treatment of (PhCH2)20 with Ac20 in a 3:1-1:1 ratio at 450-630K and 0.1-2.0 MPa for 1-200 min in a reaction zone and isolation of MeCOZCH2Ph. A mixture of (PhCH2)20 (0.3 mol) and Ac20 (0.3 mol) was heated to 573 K at 20 MPa for 20 min to give MeCOZCH2Ph with a 79% selectivity at 51% conversion of starting materials. The reaction of equimolar amts. of (PhCH2)20 and Ac20 at reflux for 25 h gave MeCOZCH2Ph with 10% selectivity after conversion of 1.4% of starting materials.

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